# Data 583 Life Expectancy (WHO)

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## Data 583 Life Expectancy - Final Report (Life Expectancy Data)

### Part 1. Introduction and Hypotheses

Life Expectancy has always been an area of interest for humanity. The dataset contains the Life Expectancy records of 193 countries between 2000-2015, together with different predictive factors. Broadly speaking, predicting variables are categorized into 4 major areas: Immmunization, Mortality, Economical, and Social, containing a total of 21 individual variables.

The primary purpose of this report is to compare and evaluate different predictive models in order to identify the most appropriate model for the dataset. In particular, we will evaluate the applicability of the core assumptions of the selected models by methods such as Normality Test, Multicollinearity Assessment, etc. This could validate or decline the adoption of certain models because the model assumptions are simply not satisfied. We are also going to verify whether the 4 predicting areas have equal significance on Life Expectancy, and whether there are adequate support evidence suggesting a strong correlation with the response variables. Finally, we will also perform and compare fitting result of selected models, in particular whether parametric models would be more suitable than non-parametric models for this dataset.

Part 2. Dataset overview

#### Varaibles Types

Variable	Unit of Measurement/Data Category	Continuous vs Discrete	Variable	Unit of Measurement/Data Category	Continuous vs Discrete
Life Expectancy	Years Old (Age)	Continuous	Total expenditure	Percentage	Continuous
Country	Nominal Data	Discrete	Percentage expenditure	Percentage	Continuous
Year	Ordinal Data	Discrete	GDP	Currency (USD)	Continuous
Status	Nominal Data	Discrete	Population	Count	Discrete
Adult Mortality	Count Data	Discrete	Income composition of resources	Percentage	Continuous
Infant deaths	Count Data	Discrete	Schooling	Mean (Years)	Continuous
Under-five deaths	Count Data	Discrete	Alcohol	Litres	Continuous
Hepatitis B	Percentage	Continuous	HIV/AIDS	Percentage	Continuous
Measles	Count Data	Discrete	$\overline{\mathrm{BMI}}$	Average BMI	Continuous
Polio	Percentage	Continuous	Thinness 1-19 years	Percentage	Continuous

Variable	Unit of Measurement/Data Category	Continuous vs Discrete	Variable	Unit of Measurement/Data Category	Continuous vs Discrete
Diphtheria	Percentage	Continuous	Thinness 5-9	Percentage	Continuous
			years		

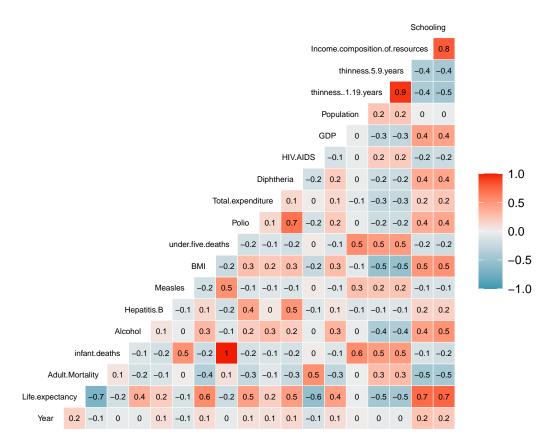
### Variables Summary and Categories

Life Expectancy is the response variable in this dataset. This represents the mean of the life expectancy (in age) in a specific country in a given year. For the data types of the predicting variables, most are percentage and count data across four major areas. The first area is Immunization Data such as Hepatitis B and Polio (immunization coverage). The second area is Mortality Data such as Adult Mortality/infant deaths (No. of deaths of Adult/infant per 1000 persons). The third area is Economical Data such as GDP/Income composition/Percentage expenditure. The fourth area is Social Data such as Schooling and Population.

To clean up and wrangle data for the subsequent analysis, NA data has been assessed. A total of 2563 NA values are found in the dataset, spreading across a few columns. These NA values are generally imputed by the respective column mean.

### Checking for Multicollinearity

As Multicollinearity can potentially affect the accuracy of regression model and we have 22 variables, a correlation study is undertaken to understand and assess the situation. A correlation plot has identified a number of correlation problems. It is found that infant deaths and under five deaths are nearly 100% correlated. The relation between the deaths rates of the two close age groups is easily interpretable. In addition, there are three heavily correlated pairs which is defined by the abs(correlation coefficient)>0.7 between the variables. They include (a) (immunization rate of) 'Polio'-vs-'Diphtheria', (b) 'income composition of resources'-vs-'Schooling', and (c) between the two thinness measures for the age groups 5-9 vs 10-19. Pairs (a) and (c) are justifiable while the relation for (b) demonstrate a relatively subtle relation. Other than that, the degree of multicollinearity is acceptable and not too worrying.



#### Variables Selection

Due to the above correlations between certain variables, in addition to a large number of variables in our model, a variable selection is conducted to remove some correlated variables and attain a simpler model via variable selection. A BIC backward step model selection method has been applied to the dataset, with the following summary:

Models	No. of Variables	AIC Score	Adj R-squared Score
Original Model	20	7642.14	0.8299
Reduced Model	12	7604.24	0.8296

The reduced model now contains the following variables: Adult.Mortality + infant.deaths + Hepatitis.B + BMI + under.five.deaths + Polio + Diphtheria + HIV.AIDS + GDP + thinness..1.19.years + Income.composition.of.resources + Schooling

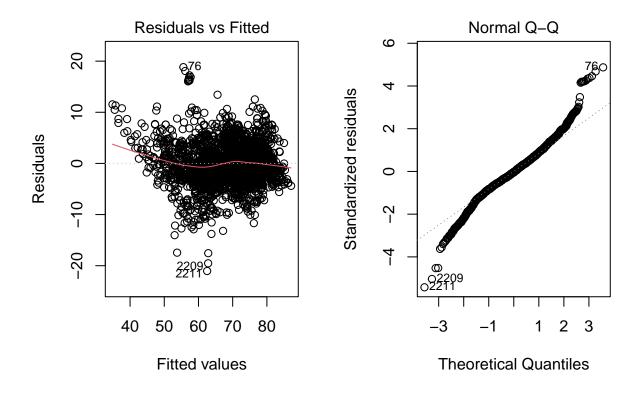
The number of independent variables are now effectively reduced to 12, together with a lower AIC score of 7604.34. Meanwhile, the adjusted R-squared score is well kept at nearly the same level as in the original model. We are satisfied with the performance of this reduced model. This reduced model will therefore be adopted as the basis for further analysis in this report.

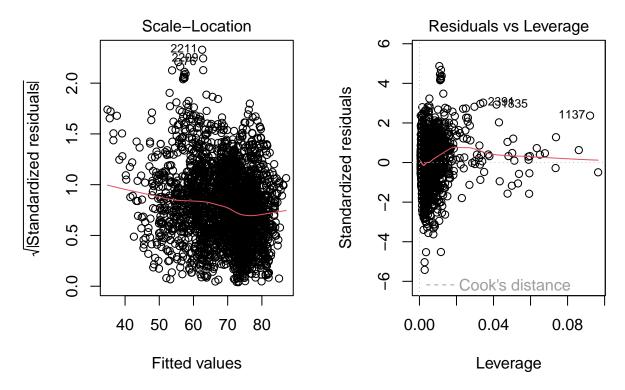
We eliminate the Status variable from the selected list of variables in the reduced model as this is a factor variable with two statuses and not continuous. We first study the effect of the model without this variable.

### Part 3. Regression Analysis

### Linear model and diagnostics

The initial model shows that we are able to explain approximately 82% of variability of our response variable using the selected predictor variables. The next step is to look at the error diagnostics from the model.

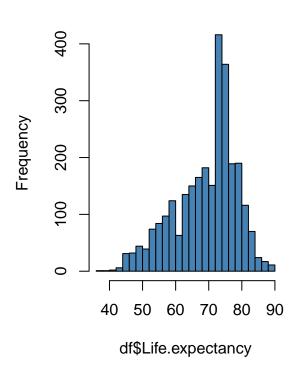


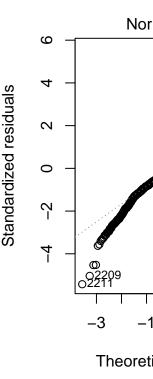


The QQ plot suggests that the model is heavy tailed and the data on both ends of the quantiles do not fit on a straight line. This is an indication that the current linear regression based model is not fitting the data well. Based on this, we undertake some additional testing to validate if the model is adequate and valid

```
#Histogram & QQPlot
par(mfrow=c(1,2))
# hist_bin_width = 4
h <- hist(df$Life.expectancy, col='steelblue', main='Life.expectancy Histogram', breaks = 35)
#not really a good "bell-shape"
# install.packages("ggpubr")
plot(lmmod2,which = 2)</pre>
```

## **Life.expectancy Histogram**





### a. Life expectancy variable distribution

```
#S #most of the data is not fall along a straight diagonal line
#qqline(df$Life.expectancy)
#Both are indicating that our predict variable Y "df$Life.expectancy" is not normally distributed
```

From the histogram, it can be noticed that the response variable life expectancy is not normally distributed. From the plot, it also seems like a bimodal distribution of life expectancy data in the dataset.

**b.** Normal distribution test for our y variable Next, we evaluate to confirm if the response variable is normally distributed using Shapiro-Wilk test. The test has a p-value that is very small and is less than 0.05, this indicates that our response variable if not normally distributed.

```
#Shapiro-Wilk Test
shapiro.test(df$Life.expectancy)

##
## Shapiro-Wilk normality test
##
## data: df$Life.expectancy
## W = 0.95676, p-value < 2.2e-16

#Finding: Since df$Life.expectancy p-value is less than .05, indicate that our y variable is not normally</pre>
```

**c.** Parametric model specification tests Another test to see if the above parametric model specification is correct.

### library(lmtest)

```
## Loading required package: zoo
##
## Attaching package: 'zoo'
   The following objects are masked from 'package:base':
##
##
       as.Date, as.Date.numeric
resettest(lmmod2)
##
##
   RESET test
##
## data: lmmod2
## RESET = 121.2, df1 = 2, df2 = 2763, p-value < 2.2e-16
d. np specification test
##
## Consistent Model Specification Test
## Parametric null model: lm(formula = Life.expectancy ~ Adult.Mortality +
##
                             infant.deaths + Hepatitis.B + BMI + under.five.deaths
##
                             + Polio + Diphtheria + HIV.AIDS + GDP +
##
                             thinness..1.19.years +
##
                             Income.composition.of.resources + Schooling, data =
##
                             df, x = TRUE, y = TRUE)
## Number of regressors: 12
## IID Bootstrap (399 replications)
## Test Statistic 'Jn': 21.17521
                                    P Value: < 2.22e-16 ***
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' 1
## Null of correct specification is rejected at the 0.1% level
```

All the diagnostic tests indicate that linear regression is not an appropriate model for the given data as assumptions for the model are violated.

#### Parametric model and assessments

As the linear model is not adequate, we move on to model this with other models that do not assume normal distribution. The models selected for the given dataset are LASSO and Neural Net with linear activation function. The following variables are selected for rest of the modeling based on correlation of the variables with the response variable and our knowledge on the domain. Here is a summary of the variable selection and our comments.

Data Categories	Vaiables
Economical Data	Total expenditure, Percentage expenditure, GDP, Income composition of resources
Social Data	Country, Status, Population, Schooling, Alcohol, BMI, Thinness 1-19 years, Thinness 5-9
	years
Mortality Data	Adult Mortality, Infant deaths, Under-five deaths

Data Categories	Vaiables
Immunization Data	Hepatitis B, Measles, Polio, HIV/AIDS, BMI, Diphtheria

	LM	LAS	SO NN	NPR	EG
Column Name	$\mathbf{Type}$				Reason of Removal
Country	(Discrete)				Since we wanted to build models for all countries
Year	(Discrete)				ordinal type data and based on domain knowledge, not consider important
Status	(Discrete)				nominal type data and based on domain knowledge, not consider important
Adult Mortality	(Discrete) X	X	X	X	1
Infant deaths	(Discrete) X	X	X	X	
Under-five deaths	(Discrete) X	X	X	X	
Hepatitis B	(ContinuouX)	X	X	X	
Measles	(Discrete)				Since it is a count and discrete type data and weak correlation with our predictor
Polio	(ContinuouX)	X	X	X	
Diphtheria	(Continuoux)	X	X	X	
Total Expenditure	(Continuous)				based on domain knowledge, not consider important
Percentage Expenditure	(Continuous)				based on domain knowledge, not consider important
GDP	(Continuou <b>X</b> )	X	X	X	importumit
Population	(Discrete)	71	71	71	no correlation with our predictor indicated by our correlation plot
Income composition of resources	(ContinuouX)	X	X	X	our correlation prov
Schooling	(ContinuouX)	X	X	X	
Alchol	(Continuous)				based on domain knowledge, not consider important
HIV/AIDS	(ContinuouX)	X	X	X	
BMI	(ContinuouX)	X	X	X	
Thinness 1-19 years	(ContinuouX)	X	X	X	
Thinness 5-9 years	(Continuous)				range already covered in 1-19 Thinness 1-19 years
status.val	(Continuous)				based on domain knowledge, not consider important

Two different supervised algorithms tried on the dataset. They do not have the constraint of a normal distribution for response variable.

First did a train and test split so we can measure the MSE and compare how each of the models are performing in terms of minimizing MSE.

Test MSE comparison for the three models

	LM	LASSO	NN
PRESS	15.93367	15.98972	22.43056

```
# #MSE comparison
# print(mselm_te1)
# print(mselas_te1)
# print(min(mse_nnet_lin))
```

Test R2 comparison for the three models

	LM	LASSO
$\mathbf{R2}$	0.829139273435324	0.829061931452822

```
# #Linear Model R2
# r2_lm<- summary(lmmodtr)$r.squared
# paste("Linear Model R2: ",r2_lm)
#
# #Lasso R2
# r2_lasso <- lasmod$dev.ratio
# paste("Lasso R2: ",r2_lasso)</pre>
```

As we compare linear model, lasso and neural net, we see that the test MSE is minimum for LASSO model. So this is a model that can be considered for the dataset.

### Nonparametric regression

The response variable shows a bimodal distribution and nonparametric regression performs better on such datasets per literature. We next try non parametric regression on the dataset.

```
library(np)
\# n \leftarrow names(df)
\# f <- as.formula(paste("df$Life.expectancy ~", paste(n[!n %in% "Life.expectancy"], collapse = " + ")))
\# model_np <- npregbw(Life.expectancy \sim Adult.Mortality + infant.deaths + Hepatitis.B + BMI + under.five.d
# model_np <- npreg(bws = model_np)</pre>
# summary(model_np)
model_np <- readRDS("model_np.rds") #PreTrained Model</pre>
summary(model np)
##
## Regression Data: 2778 training points, in 12 variable(s)
##
                 Adult.Mortality infant.deaths Hepatitis.B
                                                                   BMI
## Bandwidth(s):
                        389457535
                                                   225092161 79216285
                                         6733757
##
                 under.five.deaths
                                      Polio Diphtheria HIV.AIDS
                                                                           GDP
## Bandwidth(s):
                           95308072 5825954
                                               19248839 1.393258 167351562078
##
                 thinness..1.19.years Income.composition.of.resources Schooling
## Bandwidth(s):
                              37667667
                                                                 1071202 15165344
##
## Kernel Regression Estimator: Local-Linear
## Bandwidth Type: Fixed
## Residual standard error: 3.345092
## R-squared: 0.8722143
##
## Continuous Kernel Type: Second-Order Gaussian
## No. Continuous Explanatory Vars.: 12
```

We see that the R<sup>2</sup> is increased to 87% approximately. Done with local linear estimator and cv.aic. This is a cross validated model and help estimate the long run performance. Can we see BIC?

```
Kernel Regression Significance Test
Type I Test with IID Bootstrap (399 replications, Pivot = TRUE, joint = FALSE)
Explanatory variables tested for significance:
Adult.Mortality (1), infant.deaths (2), Hepatitis.B (3), BMI (4), under.five.deaths (5), Polio (6), Diphtheria (7), HIV.AIDS (8), GDP (9), thinness..1.19.years (10), Income.composition.of.resources (11), Schooling (12)
                  Adult.Mortality infant.deaths
389457535 6733757
Bandwidth(s):
                  Hepatitis.B BMI 225092161 79216285
                                       BMI under.five.deaths
5285 95308072
Bandwidth(s):
                    Polio Diphtheria HIV.AIDS
25954 19248839 1.393258
Bandwidth(s): 5825954
                             GDP thinness..1.19.years
Bandwidth(s): 167351562078
                                                  37667667
                  Income.composition.of.resources
Bandwidth(s):
Bandwidth(s): 15165344
Individual Significance Tests
P Value:
Adult.Mortality
infant.deaths
Hepatitis.B
                                         0.047619 *
                                            2e-16 ***
BMT
                                            2e-16 ***
under.five.deaths
Polio
Diphtheria
HIV.AIDS
GDP
                                           2e-16 ***
                                         < 2e-16 ***
thinness..1.19.years
Income.composition.of.resources
                                        < 2e-16
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Figure 1: npsigtest\_npreg result

We measure the significance of the variables for a parsimonious model. All the parameters used are significant.

Summarizing the different models and the performance assessed during the course of this project #### Add table

### Model improvements

- 1. status variable and other ordinal, nominal variables
- 2. train and test split for np, but what we have now is sufficient for measuring long run performance
- 3. multicollinearity between the variables infant.deaths and under.five.deaths. Remove one of the variables and study if there are improvements in performance
- 4. measure without imputing data

### Challenges

- 1. 30 hours for npreg
- 2. 30+ hours for model significance
- 3. null values in the dataset, columns dropped , columns imputed with mean -

### Conclusion

np and LASSO are suitable for this dataset. We find that life expectancy is....